Ser. No. 10/535,084

Response Accompanying RCE

Atty Docket 117163.00137

**AMENDMENTS TO THE CLAIMS** 

Listing of claims:

1.-3. (cancelled)

4. (currently amended) An endoprosthesis containing one or more of the elements from the

group yttrium (Y), neodymium (Nd) or zirconium (Zr), wherein the endoprosthesis is adapted to

be implanted in a vascular vessel and <u>is</u> adapted to inhibit the proliferation of human smooth

muscle cells of the vascular vessel, and wherein the formulation endoprosthesis is adapted to

liberate the one or more elements for intravascular liberation intravascularly after implantation in

a vascular vessel and the formulation endoprosthesis includes an at least very substantially

biodegradable carrier.

5.–6. (cancelled)

7. (previously presented) An endoprosthesis as set forth in claim 4, wherein the carrier is an

alloy, selected from the group consisting of magnesium, iron and tungsten alloys.

8. (withdrawn) An endoprosthesis as set forth in claim 4, wherein the carrier is a bioresorbable

polymer and one or more of the elements selected from the group consisting of Y, Nd or Zr is

embedded in the form of a powder or microparticles in the polymer.

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9. (currently amended) An endoprosthesis as set forth in claim 4, wherein the formulation

endoprosthesis contains Y in a quantitative proportion of between 3.7 and 5.5 % by weight with

respect to the total weight of the formulation endoprosthesis.

10. (withdrawn, currently amended) An endoprosthesis as set forth in claim 4, wherein the

formulation endoprosthesis contains Nd in a quantitative proportion of between 0.1 and 5% by

weight with respect to the total weight of the formulation endoprosthesis.

11. (withdrawn, currently amended) An endoprosthesis as set forth in claim 4, wherein the

formulation endoprosthesis contains Zr in a quantitative proportion of between 0.1 and 3% by

weight with respect to the total weight of the formulation endoprosthesis.

12. (currently amended) An endoprosthesis as set forth in claim 7, wherein the formulation is

<u>carrier comprises</u> a magnesium alloy and contains Y in the range of between 3.7 and 5.5%, rare

earths without Y in the range of between 1.5 and 4.4% by weight and remaining elements < 1%.

13. (currently amended) An endoprosthesis as set forth in claim 7 the formulation is carrier

comprises a magnesium alloy and contains Y in the range of between 3.7 and 5.5% by weight,

Nd in the range of between 1.8 and 2.7% by weight, and Zr in the range of between 0.2 and 1.2%

by weight.

14. (previously presented) An endoprosthesis as set forth in claim 13, wherein the magnesium

alloy is a WE43 alloy of the following formulation:

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Zirconium in an amount of 0.53 % by weight,

Yttrium in an amount of 4.1 % by weight,

Neodymium in an amount of 2.2 % by weight, and

Magnesium in an amount greater than 92.77% by weight to 93.17 % by weight.

15. (currently amended) An endoprosthesis as set forth in claim 4, wherein the formulation endoprosthesis contains Y and is so adapted that there is an yttrium concentration in the region of the human smooth muscle cells to be treated of between 200 µM and 2 mM.

16. (withdrawn, currently amended) An endoprosthesis as set forth in claim 4, wherein the formulation endoprosthesis contains Nd and is so adapted that there is a neodymium concentration in the region of the smooth muscle cells to be treated of between 600 µM and 2 mM, in particular between 800 µM and 1 mM.

17. (withdrawn, currently amended) An endoprosthesis as set forth in claim 4, wherein the formulation endoprosthesis contains Zr and is so adapted that there is a zirconium concentration in the region of the smooth muscle cells to be treated of between 200  $\mu$ M and 2 mM, in particular between 200  $\mu$ M and 1 mM.

18. (withdrawn, currently amended) An endoprosthesis as set forth in claim 4, wherein the formulation endoprosthesis contains Y, Nd and Zr and is so adapted that there is an yttrium concentration of between 350 and 550 µM, a neodymium concentration of between 100 and 200

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 $\mu M$  and a zirconium concentration of between 10 and 30  $\mu M$  in the region of the smooth muscle cells to be treated.

19.-25. (cancelled)

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